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Epidemic potential of an emerging vector borne disease in a marginal environment: Schmallenberg in Scotland

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During 2011 Schmallenberg virus (SBV) presented as a novel disease of cattle and sheep that had apparently spread through northern Europe over a relatively short period of time, but has yet to infect Scotland. This paper describes the development of a model of SBV spread applied to Scotland in the event of an incursion. This model shows that SBV spread is very sensitive to the temperature, with relatively little spread and few reproductive losses predicted in years with average temperatures but extensive spread (>1 million animals infected) and substantial reproductive losses in the hottest years. These results indicate that it is possible for SBV to spread in Scotland, however spread is limited by climatic conditions and the timing of introduction. Further results show that the transmission kernel shape and extrinsic incubation period parameter have a non-linear effect on disease transmission, so a greater understanding of the SBV transmission parameters is required.

Reports of short-term pyrexia, diarrhoea and reduced milk production in dairy herds in Western Germany in August 2011 prompted identification of a new emerging disease, that became known as Schmallenberg virus^{1,2} (SBV). An Orthobunyavirus of the Simbu serogroup, SBV is closely related to Akabane and Shamonda viruses. Similar to Akabane virus, SBV infection can cause clinical disease in cattle, sheep and goats, characterised by pyrexia, reduced milk production, abortions and congenital malformations among offspring whose mothers are infected during a particular period of pregnancy^{1,2}. Aside from congenital infection, the main transmission route is via arthropod vectors (principally *Culicoides* spp)^{1,3–6}. During late 2011 and early 2012 cases were identified in a number of northern European countries⁷ including 276 farms in England and the Channel Islands where SBV was identified in aborted or malformed calves or lambs (as of 26th July 2012)⁸.

As a newly identified pathogen, the epidemiology of SBV is not yet fully known. However, information available so far on the clinical picture and transmission routes suggest it is sufficiently similar to Akabane virus to use knowledge of Akabane to assist our understanding of SBV⁹. Furthermore, recent studies of the epidemiology of another *Culicoides*-borne arbovirus, Bluetongue virus (BTV), can be used to draw inference for the epidemiology of SBV within vector populations. Previous studies have suggested that the potential for spread of arboviruses in Europe may be driven by climate change^{10–12} and such changes, specifically temperature increases, may determine the potential for spread of SBV and other arboviruses in Scotland.

Previous studies^{13–15} have characterised the *Culicoides* populations in Scotland which include a number of species, principally *C. pulicaris* and *C. obsoletus* that feed on wildlife and livestock species, and one species *C. impunctatus* that also feeds on humans. These species have very different ecologies^{13,15} with *C. impunctatus* dominating in bog and heathland habitats whilst other *C. pulicaris* complex species and *C. obsoletus* complex species are primarily associated with pastoral livestock farming.

Some information is available on vectoral capacity of these species for BTV that also has relevance for SBV transmission. Relative to *C. obsoletus* and *C. pulicaris* complex species, *C. impunctatus* is thought to be an inefficient vector species, due to its wide host preference including humans¹⁶, lower ability to replicate BTV in the laboratory than other Palaearctic species (0.4% versus 0.5–13% in *C. obsoletus* and *C. pulicaris* complex species^{17,18}) and autogeny, where the female may lay its first batch of eggs without taking a bloodmeal¹⁹. However, its high population densities in Scotland and northern England (landing rates of 10–635 midges/min on an exposed human arm¹⁷) may compensate for these characteristics and allow *C. impunctatus* to transmit BTV and potentially SBV.



At the time of writing there have been no confirmed cases of animals being infected in Scotland. One seropositive animal has been found, that is believed to have been infected prior to import from southern England. However, due to the proximity to England and the large abundance of *Culicoides* vectors throughout Scotland^{13,15} there remains a risk of introduction of SBV into the Scottish vector and livestock population. Assessment of the likely extent of spread and ramifications in Scotland would aid control and deployment of resources following introduction. Therefore, the aims of this paper are as follows:

1. To investigate the likely extent and impact of an SBV epidemic in Scotland.
2. To explore the effect of temperature on the risk of SBV spread in Scotland and the potential effect of climatic warming on SBV transmission.
3. To explore the importance of the parameters of SBV transmission, in particular the parameters relating to the vector.

Results

A stochastic simulation model was used to investigate the likely transmission of SBV in Scotland. The model incorporates the transmission of disease from host to vector and from vector to host following the introduction of disease (seeding) on between 1 and 10 farms. Transmission between and within individual farms is determined by a transmission kernel that describes the vector dispersal potential. The distribution of the number of sheep and cattle infected under the baseline model is shown in Figure 1.

Under the baseline scenario using the mean temperature the resulting number of animals infected is relatively low (Table 1, Figure 1). There is a small epidemic if disease is introduced on day 60, with around 14 ewes with potential reproductive losses (Table 1, Figure 1). There is little disease spread if there is introduction earlier or later than day 60.

Under the extreme case scenario using the maximum observed temperature (mean difference between mean and maximum temperature is 1.96°C), there is substantial disease spread (Table 1, Figure 2).

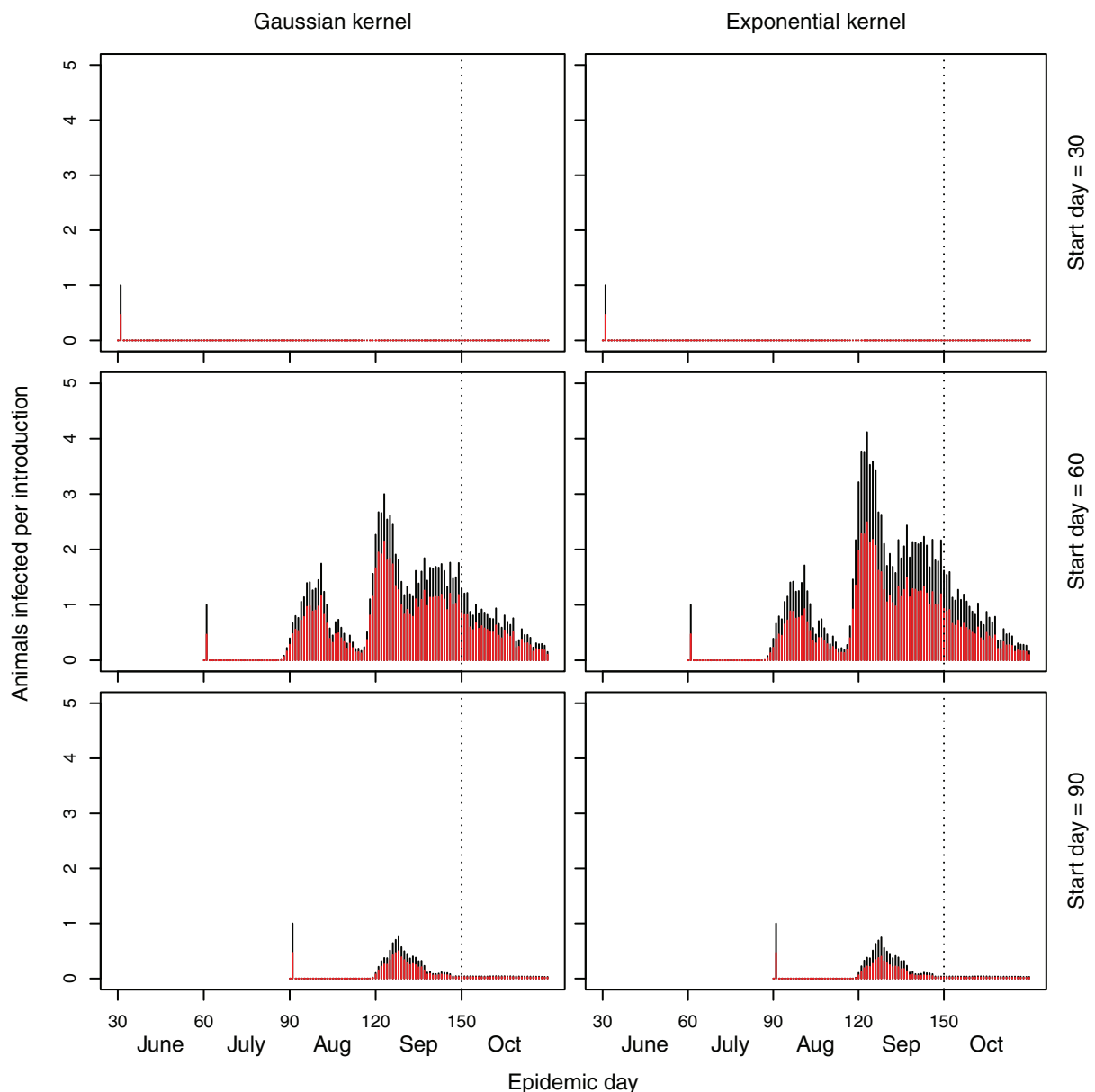


Figure 1 | Stacked barplots of the number of sheep (red bars) and cattle (black bars) infected on each day of the simulated epidemic under the baseline implementation in which the mean temperature is used. The dashed line represents the start of the period in which in-lamb ewes may be at risk of reproductive losses.

**Table 1 | Summary of results from the different model implementations in this study**

Baseline model—mean temperature						
Start day	Kernel	Proportion spread	mean (sd) farms affected	mean (sd) cattle infected	mean (sd) sheep infected	mean (sd) ewes at risk*
30	Gaus	0.009	5.51 (2.89)	2.93 (1.92)	2.59 (1.80)	0.00 (0.01)
	Exp	0.01	5.52 (2.90)	2.94 (1.95)	2.59 (1.80)	0.00 (0.01)
60	Gaus	0.983	368 (242)	172 (161)	379 (351)	13.0 (18.2)
	Exp	0.981	300 (203)	275 (295)	401 (427)	12.8 (20.2)
90	Gaus	0.982	52.0 (47.4)	21.2 (39.1)	38.3 (67.5)	0.73 (7.09)
	Exp	0.980	46.3 (31.3)	26.9 (52.0)	31.9 (51.4)	0.57 (5.44)
120	Gaus	0	5.49 (2.87)	2.93 (1.93)	2.56 (1.77)	0.00 (0)
	Exp	0	5.49 (2.86)	2.94 (1.94)	2.55 (1.77)	0.00 (0)
Extreme case scenario—maximum temperature						
30	Gaus	0.980	5603 (1936)	331127 (185643)	919183 (587984)	29640 (17277)
	Exp	0.982	2848 (1333)	246903 (131765)	513767 (306481)	14470 (9959)
60	Gaus	1.000	6106 (1548)	308016 (130705)	783905 (362910)	48410 (24301)
	Exp	1.000	3500 (1411)	271761 (118586)	526484 (246401)	24317 (12130)
90	Gaus	1.000	1415 (544)	1855 (1740)	3972 (3414)	151 (255)
	Exp	1.000	964 (450)	3162 (2737)	4298 (3372)	141 (278)
120	Gaus	0.949	95.5 (63.4)	36.8 (32.5)	70.7 (59.9)	11.5 (10.0)
	Exp	0.948	81.4 (50.9)	47.8 (41.0)	59.2 (48.5)	9.60 (8.08)
Mean temperature—EIP minimum incubation temperature – 1°C						
30	Gaus	0.897	1217 (1149)	3675 (5603)	8796 (13700)	621 (929)
	Exp	0.898	718 (654)	7883 (11346)	12483 (18092)	667 (966)
60	Gaus	0.999	3109 (1132)	29087 (22563)	62522 (48571)	7181 (5775)
	Exp	0.999	1876 (842)	57183 (41815)	83823 (63229)	8981 (6596)
90	Gaus	0.998	355 (223)	204 (340)	397 (576)	43.0 (57.5)
	Exp	0.998	307 (197)	402 (557)	476 (560)	59.6 (66.8)
120	Gaus	0.302	10.8 (11.8)	5.07 (5.42)	6.07 (7.79)	0.59 (1.24)
	Exp	0.307	9.92 (9.55)	5.98 (7.07)	5.09 (6.00)	0.43 (0.93)
Mean temperature—EIP Incubation rate + 25%						
30	Gaus	0.036	5.72 (4.31)	2.99 (2.10)	2.75 (3.22)	0.00 (0.02)
	Exp	0.034	5.82 (6.03)	3.04 (3.02)	2.90 (6.17)	0.00 (0.02)
60	Gaus	0.989	973 (525)	837 (732)	1912 (1675)	25.6 (44.3)
	Exp	0.989	661 (385)	1355 (1275)	1982 (1817)	25.5 (42.6)
90	Gaus	0.987	90.9 (72.6)	41.3 (122)	76.4 (202)	1.64 (19.6)
	Exp	0.987	78.1 (49.5)	54.1 (175)	63.8 (159)	1.27 (14.5)
120	Gaus	0	5.49 (2.87)	2.93 (1.92)	2.57 (1.78)	0
	Exp	0	5.49 (2.87)	2.93 (1.92)	2.57 (1.79)	0

*Ewes at risk of reproductive losses.

The key determinant of the size of epidemic is the date of introduction (Table 1, Figure 2). The greatest number of infected animals follows a disease introduction on day 60 (30th June) as this period corresponds with the highest temperatures and therefore the highest potential for spread. Infection on day 90 results in only a small number of infected animals (Figure 2). The epidemic spreads in distinct waves (Figure 1) that correspond with the Extrinsic Incubation Period (EIP). The number of sheep that are at risk of reproductive losses is considerably higher with an introduction on day 60 compared to introduction on day 30. The proportion of infected cattle that are at risk of reproductive losses is not shown because: 1) the window at which cattle are at risk is much wider (between days 62nd and 173rd day of pregnancy²⁰) and 2) the window over which cattle are serviced (mated) is widely distributed through the year. Therefore many of the cattle infected may be at risk of reproductive losses, but these factors make the precise impact harder to predict than they are in sheep.

A further key determinant of the extent of the size of the epidemic under the maximum temperature scenario is the number of seedings. For an epidemic started on day 60, the number of infected farms increases by around 360 for each additional disease introduction

(Figure 3; linear regression estimate 374.7 for the Gaussian kernel and 344.8 for the exponential kernel, both $p < 0.001$).

Under the baseline (mean temperature) scenario, the model is slightly sensitive to raising the incubation rate of the EIP by 25% with slightly larger epidemics with disease introduced on day 60 (Table 1, Figure 4). However, the effect of lowering the minimum temperature of the EIP by 1°C has a large effect on the size of the epidemic, with an increase of around 200 times on the number of animals infected compared to the baseline scenario (Table 1, Figure 4).

The distribution of parish level risk of transmission illustrated by the expected number of infectious vectors resulting from an infected animal on day 60 under the extreme case maximum temperature scenario is shown by Figure 5. There is a distinct concentration of higher risk in the south-west with some patches of higher risk in northerly areas.

Discussion

This paper has described the development and results of a model to combine both within farm and between farm spread of SBV using knowledge already acquired of SBV and Akabane virus epidemiology

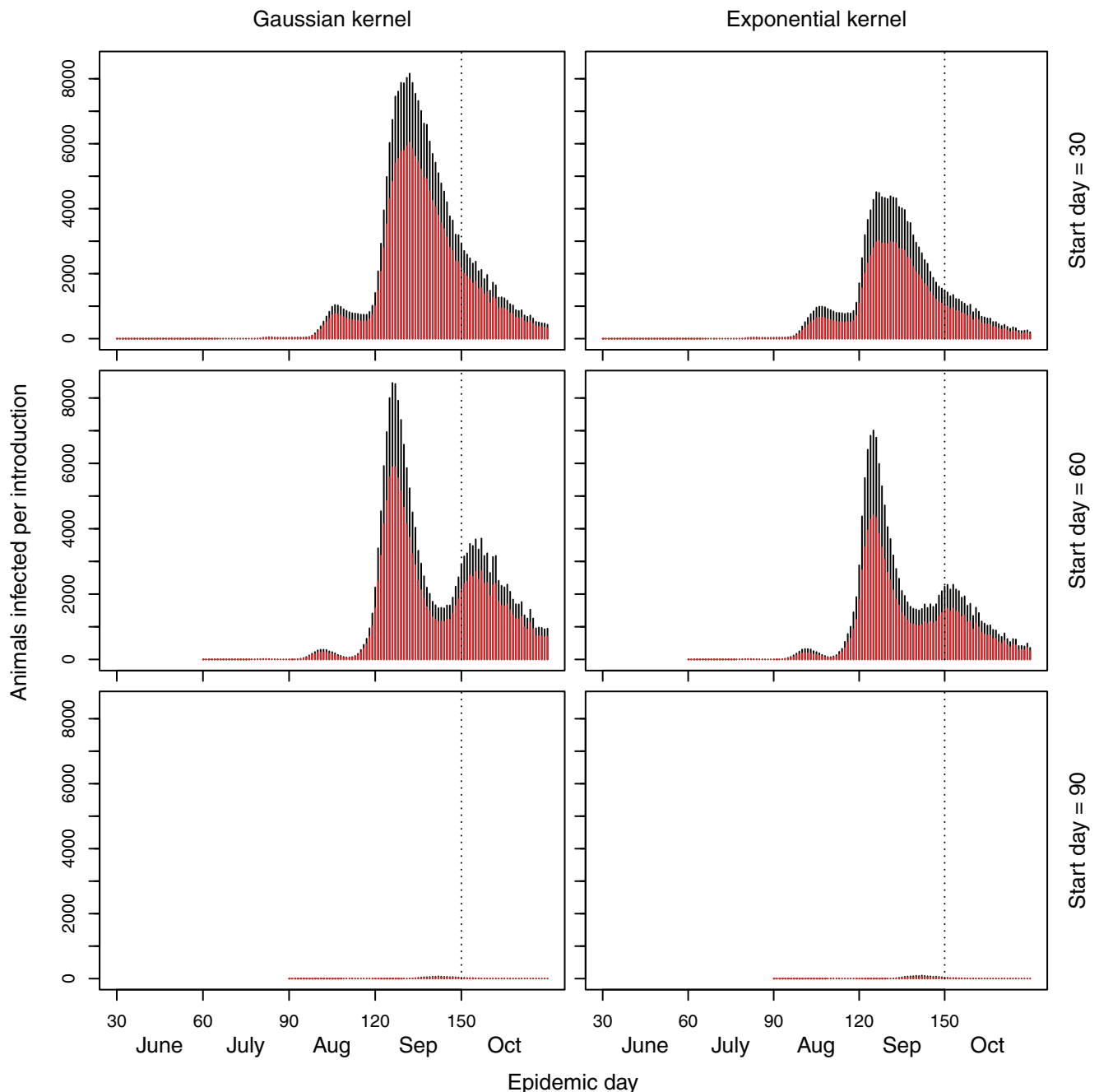


Figure 2 | Stacked barplots of the number of sheep (red bars) and cattle (black bars) infected on each day of the simulated epidemic when the maximum temperature is used. The broken line represents start of the period in which in-lamb ewes may be at risk of reproductive losses. Note the different scale compared to Figure 1.

then applied to the Scottish context using studies of *Culicoides* ecology and understanding of BTV. The model simulates introduction of the virus through windborne spread from England or Ireland or introduction through the movement of infected animals into Scotland. The findings of this paper support the findings of similar studies of BTV that shows that it would be possible for *Culicoides* borne disease to spread in Scotland^{21,22}.

Under the extreme case scenario, assuming of the maximum observed temperatures, the extent of disease spread depends heavily upon the time of introduction during the vector season. Introduction on day 30 or 60 result in similar sized epidemics, however, the epidemics are around 100 times smaller following an introduction on day 90 (Figure 2). Introduction on day 120 results in very small numbers of secondary cases (Table 1). Under the baseline, mean

temperature scenario, large epidemics only occur with introduction on day 60. These differences associated with the timing of introduction are the result of a number of factors, principally the remaining period of vector activity, bi-modal distribution of *Culicoides* abundance, temperature at the time of introduction and the remaining duration of the season of vector activity. Assuming that the disease is introduced through England or Ireland, monitoring the disease spread in these countries could be used to predict the likely arrival time in Scotland and therefore the extent of spread after arrival. Multiple seedings are used here to initiate the epidemic because multiple farms may be at risk from such wind borne vector introductions²³.

The model results suggest that given the maximum temperature conditions, the impacts of SBV in terms of reproductive losses among

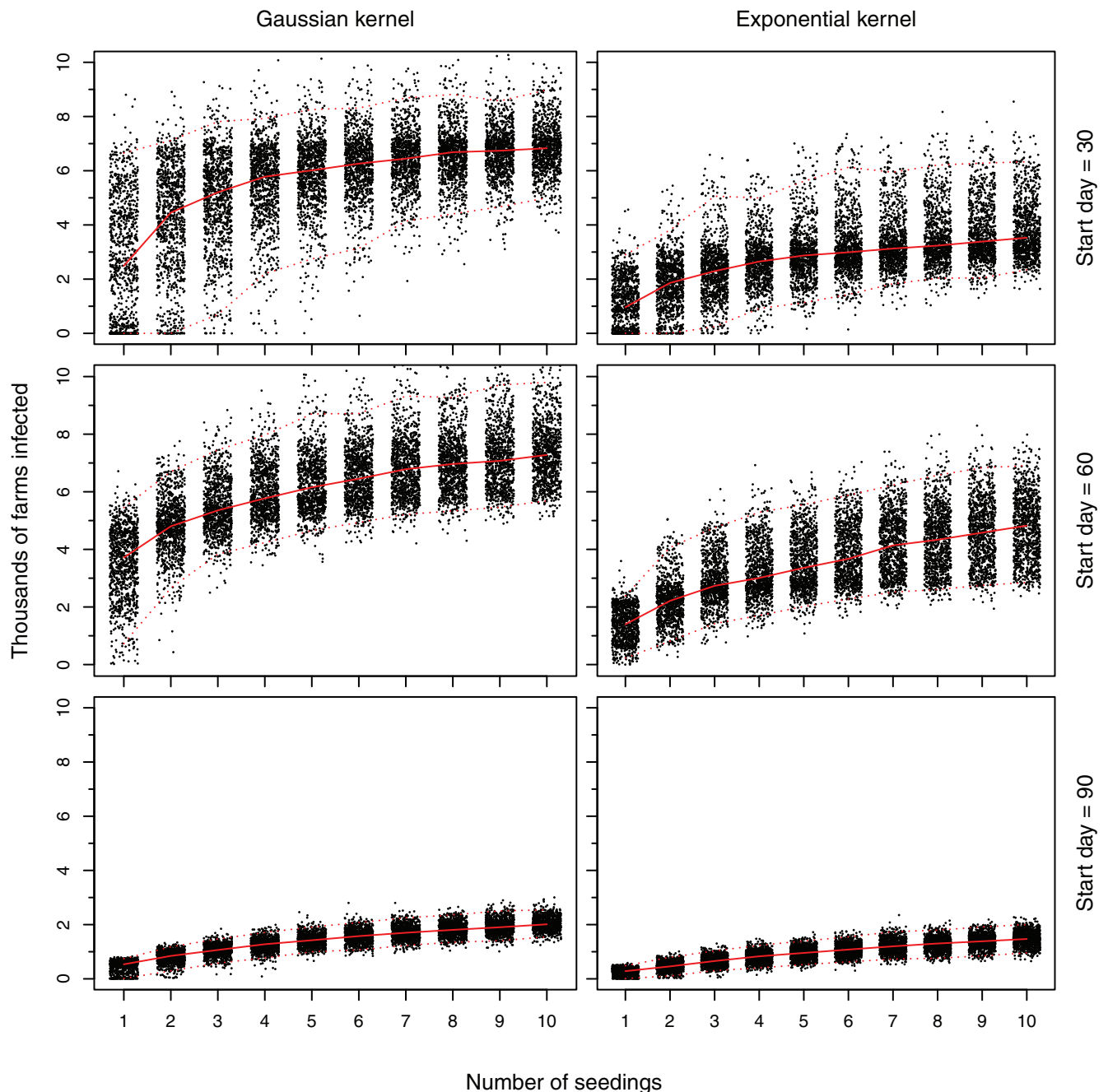


Figure 3 | The number of farms infected against the number of disease introductions under the extreme case maximum temperature scenario. The solid red line represents the median and the broken red lines the 2.5 and 97.5 percentiles. The points have been jittered to ease interpretation.

sheep will be large. This is in agreement with reports from England and elsewhere in Europe^{20,24}, where widespread birth malformations among sheep have been reported. Losses in Scotland could be reduced by delaying tupping by 15 days, thus avoiding a substantial proportion of the at-risk window (Figure 2). These results are based on the assumption that one third of ewes will be tupped during October for lambing during February, which may be inaccurate. However, the earliest tupping flocks are likely to be in the lowland, southern areas that are also the highest risk areas for SBV (Figure 5). Furthermore, this model assumed that all pregnant animals infected during the at risk period are at risk of reproductive losses, but the true proportion of these animals would go on to suffer abortions or birth malformations is currently unknown, so the observed number of reproductive losses in an epidemic may be lower. In addition to losses among sheep, a large number of cattle will become infected and this

could manifest clinically in drops in milk production or calf birth malformations and reproductive losses.

Under the baseline scenario, the size of the epidemic was exponentially smaller than the maximum temperature scenario, with around one thousandth of the numbers of animals infected (Table 1, Figure 4). However, these analyses show that even during an average year SBV does present a risk to livestock in Scotland. If average temperatures and frequency of warm periods in Scotland increase as a result of global warming then the risk of vector borne disease spread in Scotland and elsewhere in Europe will continue to increase.

At the time of writing there was no published literature on the EIP of SBV infection within *Culicoides* vectors. As a result, the EIP had to be taken from studies of BTV and this parameter may not be an accurate reflection of the infection process of SBV within

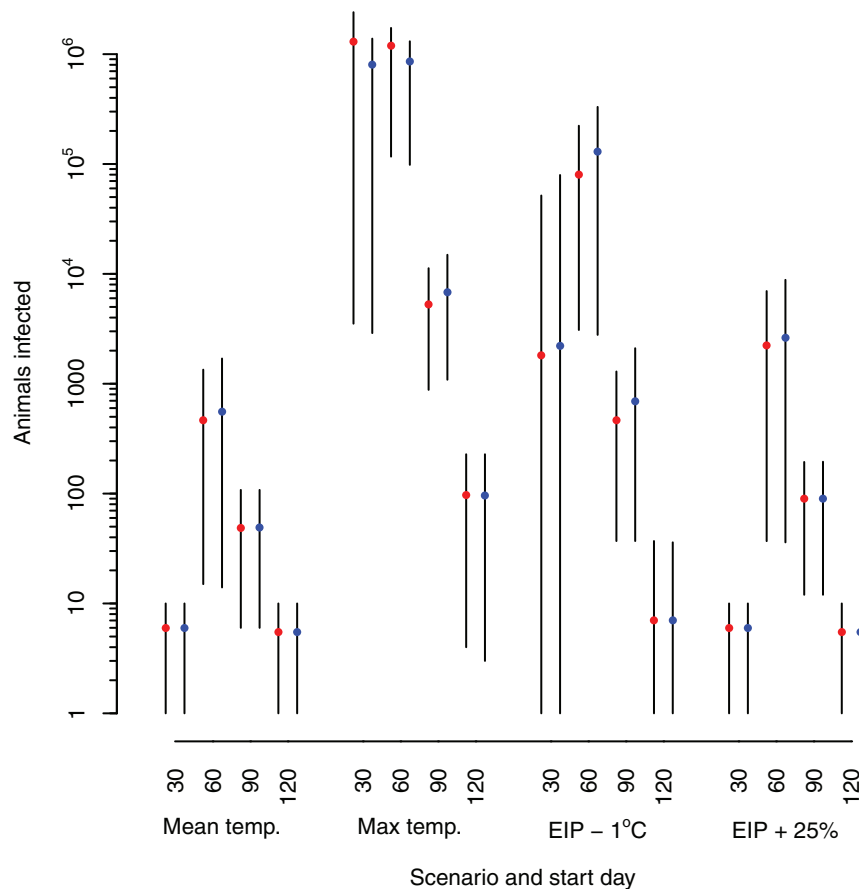


Figure 4 | Comparison of the numbers of animals infected under the scenarios in Table 1. Red points correspond to the exponential kernel, blue points to the Gaussian kernel. The black lines represent the range from the 10th to the 90th percentiles.

Culicoides. As a result we explored alternative parameters for the EIP that would result in more efficient disease transmission. Reducing the interval between infection cycles by increasing the EIP incubation rate by 25% results in around 4 times more animals infected (Table 1, Figure 4). Alternatively reducing the baseline temperature for virus incubation results in a substantially larger epidemic, by both increasing the numbers of areas with appropriate conditions for spread and increasing the rate of spread by lowering the temperature is possible (Table 1, Figure 4). The EIP incubation rate 25% faster and the baseline temperature 1°C cooler are within the ranges of values and their 95% credible limits of parameters for the EIP of BTV and African Horse Sickness identified in a recent review by Carpenter et al²⁵, so can be considered as credible values for SBV. The sensitivity of the model to these parameters highlights the need to better understand the parameters for SBV transmission if we are to accurately predict the spread of the disease.

Rather than separating within farm from between farm spread, this model incorporated both into a single transmission kernel to reflect the blood feeding behaviour of the vector. The kernel did not directly allow for wind-borne transmission that could be incorporated using an irregular shaped kernel. However, it has been demonstrated elsewhere that the role of wind in spreading vector borne disease over land is relatively minor²⁶ and that the relatively unpredictable nature of the wind speed and direction over complex topography makes this difficult to use in a predictive model. Furthermore, under the maximum temperature scenario the Gaussian kernel resulted in more animals on more farms being infected. This is because the Gaussian kernel allows the virus to spread more rapidly to new areas, whereas the exponential kernel limits transmission and ensures that locally the epidemic burns-out faster. However the

reverse is true when the mean temperature is used. This is because under the lower temperatures there are fewer areas suitable for spread. The tighter exponential kernel is more effective in ensuring that infection remains within these areas, whereas the wider Gaussian kernel allows more opportunity for the virus to escape these risk areas to areas that are less favourable for spread and conditions are less likely to lead to further transmission.

In conclusion, the climatic conditions in Scotland are at the lower end of the scale that is suitable for SBV transmission, but are still adequate for the disease to spread. However, given warmer than average (but still feasible) temperatures SBV could spread very quickly within Scotland with significant impacts in both the sheep and cattle populations. This is upon the assumption that SBV will spread with similar parameters as BTV and is highly dependant upon the nature of vector movement with respect to a transmission kernel. Furthermore, how the disease spreads within Scotland is very dependant upon the precise nature of the EIP of SBV.

Methods

The model used in this analysis is a stochastic simulation model comprising two main compartments: a) the transmission from livestock host to vector; and b) the transmission from vector to host. Different introduction scenarios are simulated by seeding of infection on to farms to represent importation of infected animals or windborne virus spread.

A number of assumptions have been made in this model:

1. Movements of exposed or infectious animals are not explicitly considered. A movement of an animal during its short latent or infectious period could spread disease to different areas of the country but was not considered here due to the fluidity of the movement network^{27–30}. In these analyses we use one kernel that implicitly includes movements³¹ and one that does not²⁶.
2. The range of dispersal of the vector can be modelled using a kernel. Vector dispersal is influenced by many factors including weather^{26,32}. However,

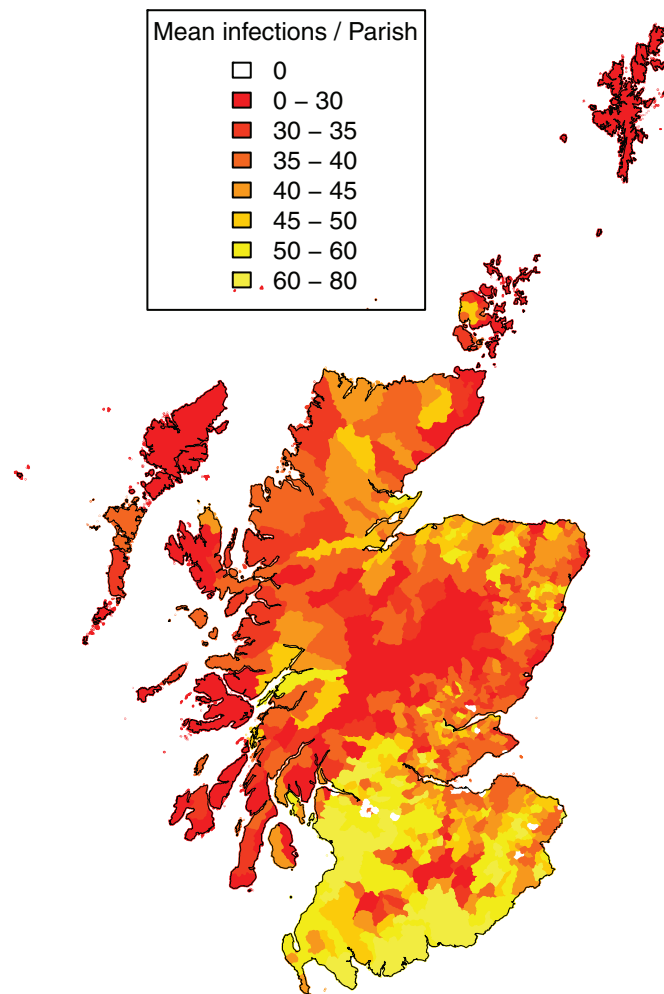


Figure 5 | The expected number of infectious bites (σ_{it}) resulting from an infected host on the 30th June (day 60) under the extreme case maximum temperature scenario. This is shown as the parish level mean.

3. The animal is equally infectious on each day of its infectious period.
4. Once infected an animal will recover with full immunity and will not be susceptible to further infection.
5. As with other studies^{20,31,34} *Culicoides* vectors are assumed to be equally likely to feed on a cow as a sheep. Studies on *C. imicola*, indicate that the vector has a preference for feeding on cattle rather than sheep. However, *C. imicola* is not a species native to Scotland. The only study of species found in Scotland show that insects have a preference for horses over cattle and sheep, but insufficient insects were trapped on cattle and sheep to make further inferences³⁵.
6. The attractiveness of a farm for vector feeding is based on the number of livestock on the farm and is determined by distance and the number of livestock.
7. Ewes are at risk of reproductive losses when infected between days 28–56 of pregnancy²⁰ and most sheep in Scotland are tupped (mated) between September and November for lambing between February and April. Therefore, in this model, any in-lamb ewe that is infected after day 150 (during October) is at risk of reproductive losses including birth malformations and abortions. These ewes are just those that were tupped during September: we assume that one third of breeding ewes are tupped during September and the other two thirds during October and November. The proportion of the sheep population that is maintained as breeding stock is 51% on the Scottish June agricultural census. The corresponding at-risk period for cattle may be as wide as 62–173 days into pregnancy²⁰ and cattle births are distributed more evenly through the year relative to sheep births.
8. Scotland can be regarded in isolation. During an epidemic involving southern Scotland there is likely to be some transmission with farms in northern England. However, as this is likely to be a two-way exchange we consider that this would have minimal effect on the epidemic.
9. No disease control measures will be employed.
10. The EIP for SBV will be the same as for BTV. This is explored in sensitivity analysis.

Model description (a) – animal to vector transmission. For an infectious farm i , at time t the number of vectors that will feed on an infectious animal and survive sufficient time to lay an infectious bite is defined by a daily rate σ_{it} . The actual number of infectious bites is taken from a Poisson distribution ($\text{Pois}(\sigma_{it})$). σ_{it} is defined as:

$$\sigma_{it} = n_{v(t)} i_{v(t)}^* \sigma_v \sigma_{a(i)} \quad (1)$$

where

$$n_{v(t)} = n_v v_t v_i \quad (2)$$

describes the number of vectors feeding on an infected animal with n_v as the expected number of vectors (set to a fixed value of 2500³⁶), v_t adjusts for the expected number of vectors at time t , following a bimodal distribution based upon Searle et al¹⁴:

Table 2 | The parameters used in this model

Parameter	Description	Derivation	Values
σ_{it}	Rate describing the number of infected flies from a single infected animal on a single day. Inserted into a random Poisson distribution.		≥ 0
n_v	Expected number of vector bites per day	After Gerry et al ³⁶ and consistent with studies from Scotland Searle et al ¹⁴ , Blackwell et al ¹⁵ .	2500
v_i	Spatial adjustment to n_v based upon landscape suitability and <i>Culicoides</i> species competence.	After Purse et al ¹³ .	$0.5 \leq v_i \leq 1$
L^p	Proportion of 1 km buffer identified as pasture	CORINE landcover 2000 ⁴⁰	≤ 1
L^h	Proportion of 1 km buffer identified as heathland	CORINE landcover 2000 ⁴⁰	≤ 1
v_t	Temporal adjustment to n_v based upon temporal peaks in <i>Culicoides</i> abundance	After Searle et al ¹⁴ .	0.25, 1
T	Temperature.	See supplementary material	9.3 – 19°C
v_m	Temperature dependant vector mortality rate.	After Gerry et al ³⁷	< 1
v_e	Extrinsic (within vector) incubation period	After Carpenter et al ²⁵ work on BTV	
v_b	Time interval between bites	After Mullens et al ³⁸ work on BTV	
σ_v	Probability of transmission between host and vector.	After Veronesi et al ³	0.19
n_L	Number of susceptible livestock.	From the Scottish June agricultural census.	
σ_T	Probability of transmission from vector to host	Derived from Baylis et al ³⁹	0.77
latent period	Period between infection and viraemia within the animal	Approximation from experimental infections of 3 cattle ¹ .	2 days
Infectious period	Period of infectiousness of an animal	Approximation from experimental infections of 3 cattle ¹ .	4 days

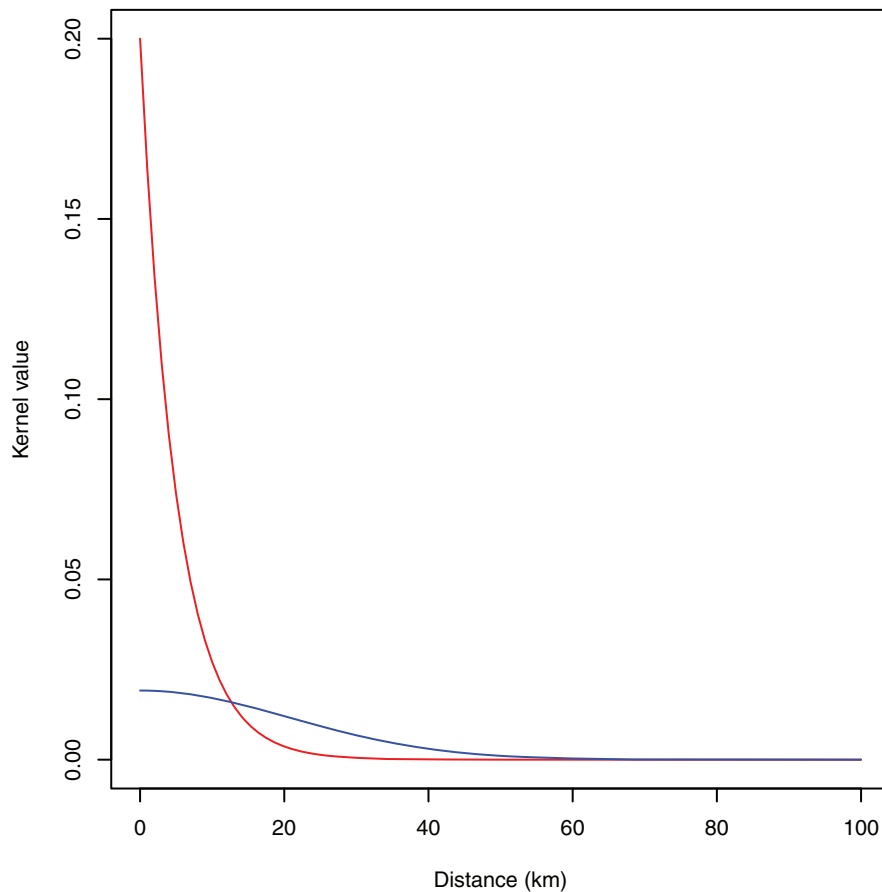


Figure 6 | The shape of the exponential kernel (red line) and Gaussian kernel (blue line).

$v_t = 0.25$ when $t < 30$ (31st May) or $t > 140$ (September 18th) and t is between 65 (5th July) and 100 (9th August) days
 $v_t = 1$ otherwise.

v_i is a term that describes the likely vector abundance given the landcover in a 1 km buffer around i , with the more competent species associated with pasture and less competent species associated with heathland:

$$v_i = 0.5 + 0.5L_i^p + 0.25L_i^h \quad (3)$$

where L_i^p and L_i^h are the proportion of land within the 1 km buffer classed as pasture and heathland respectively.

$\sigma_{a(t)}$ is the number of infectious animals on i at time t .

σ_v is the probability of transmission from animal host to vector and is equal to 0.19, based upon experimental infections³.

$\gamma_{v(t)}$ describes the probability of a midge surviving sufficient time to lay one or more infectious bites and is dependent upon three parameters: the vector mortality rate³⁷ (v_m), the EIP (V_e) and the interval between blood meals (v_b).

$$v_m = 0.009e^{0.16T} \quad (4)$$

where T is the temperature and the daily survival probability is e^{-v_m} .

The time to the vector being infectious is described by the EIP which has been parameterized for BTV, since no value is available for SBV (derived from Carpenter et al¹⁷):

$$v_e = (\max(0, 0.019(T - 13.3)^{-1})) \quad (5)$$

Following the vector becoming infectious, the time until the first potentially infectious blood meal is taken as half the interval between blood meals³⁸ where:

$$v_b = \left(0.0002T(T - 3.7)(41.9 - T)^{\frac{1}{27}}\right)^{-1} \quad (6)$$

In order to account for a vector taking more than one potentially infectious meal, the infectious bite rate of a vector is given by:

$$\gamma_{v(t)} = \sum_{i=0}^{i \leq 10} (e^{-v_m})^{v_e + 0.5v_b + v_b i} \quad (7)$$

This assumes that the midge will not survive to lay more than 10 infectious bites (although the probability of surviving more than two bites being very small and is almost negligible).

Model description (b) – vector to animal transmission. An infected vector from farm i will infect an animal on farm j (i can be the same as j) with probability defined by:

$$p_{ji} = p_j^{(A)} \kappa(\chi_{ij}) \lambda_i \quad (8)$$

where $\kappa(\chi_{ij})$ is a spatial transmission kernel based upon the Euclidean distance between farms i and j (χ_{jk}) and describes the distances a vector will travel between infection and the laying of infectious bites. $p_j^{(A)}$ describes the attractiveness of a farm to a vector, and is equal to the number of susceptible hosts ($n_j^{(L)}$) on holding j .

λ_i is a scaling parameter that ensures that any potentially infectious bite is laid only once:

$$\lambda_i = \frac{1}{\sum_{j \in \mathbb{R}} (p_j^{(A)} \kappa(\chi_{ij}))} \quad (9)$$

The probability of successful transmission from vector to host is given by $\sigma T = 0.77$ ³⁹ and following infection the animal is latently infected (non-infectious) for two days and infectious for four days¹.

The model parameters are summarised in Table 2.

Model implementation. The model assumes a potential vector activity season from May to October (inclusive) which is within the duration of vector activity observed in studies in Scotland^{14,15}. The model introduces infection on different start dates from the 1st June (being the first month at which temperatures are suitable for within vector incubation) with 30 day intervals between start days (up to start day = 120 (29th August)).

The model is initiated by introducing disease on certain 'seed' farms. The seeds are assumed to be introductions from England, so a farm is more likely to be a seed if it is further south (described in the supplementary information) with seeds sampled at random (with replacement, so it is possible for a seed farm to become infected twice). The seed is infected by a single random animal becoming infected.



Two shapes of transmission kernel $\kappa(x_{ij})$ are considered; a Gaussian kernel based upon Szmarađ et al.³¹, described by $\frac{\alpha}{\sqrt{\pi}} e^{-\alpha^2 x^2}$ where $\alpha = 0.034$ and an exponential kernel $0.2e^{-0.2x}$ based upon Sedda et al.²⁶ (Figure 6). For each combination of parameters (two kernel shapes, 4 start days, 1 to 10 seeds) 1000 iterations were run resulting in 80000 model runs.

Data. The unique parish-holding (PH) identifiers were taken from the Scottish June agricultural census for 2011. This comprised 52,543 holdings of which the 20,877 with at least one head of cattle or one sheep were selected. Coordinates for the holdings were identified from Animal Movement Licensing Scheme (AMLS) location data or British Cattle Movement Scheme (BCMS) Cattle Tracing System (CTS) Postcode Address Field (PAF) data. Landcover variables were derived from the CORINE 2000 Landcover map⁴⁰.

Temperature data were extrapolated for each farm from the UK Met Office UK CIP archive of temperature data interpolated and gridded to 5 km² cells between 1990 and 2006⁴¹. These data were processed using the methodology described in Supplementary material to extract the mean monthly temperature for each cell during the period in question.

Sensitivity analysis. To explore the sensitivity of the model to certain parameters sensitivity analysis was conducted by altering the parameter and rerunning the model with the new values and comparing these results to those from the baseline (mean temperature) scenario. Analysis was conducted to explore the effects of using the maximum mean monthly temperature of each grid cell over the period of the data (1990–2006). The parameter for EIP is taken from studies of BTV and the parameter for SBV may have a different minimum incubation temperature or a different virus replication rate. Thus an equation with a minimum temperature 1°C cooler:

$$v_e = (\max(0, 0.019(T - 12.3))^{-1}) \quad (10)$$

and a virus replication rate 25% faster:

$$v_e = (\max(0, 0.02375(T - 13.3))^{-1}) \quad (11)$$

were used in sensitivity analysis.

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Author contributions

P.R.B., B.M.deC.B. and I.G.H. conceived and designed the study, P.R.B. performed the analysis and drafted the manuscript, K.R.S. and B.V.P. contributed to aspects of arbovirus epidemiology and HKA contributed to veterinary aspects of SBV. All authors have read, made changes to and approved the manuscript.

Additional information

Supplementary information accompanies this paper at <http://www.nature.com/scientificreports>



Competing financial interests: The authors declare no competing financial interests.

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